

(FILE 'HOME' ENTERED AT 11:44:10 ON 25 JUL 2002)

FILE 'MEDLINE, CAPLUS' ENTERED AT 11:44:25 ON 25 JUL 2002

L1 360 S SEROTONIN TRANSPORTER GENE
L2 0 S L1 AND (INFECTIOUS DISEASE? OR FLU OR INFLUENZA OR PNEUMONIA
L3 73 S L1 AND DISEASE?
L4 58 DUP REM L3 (15 DUPLICATES REMOVED)
L5 58 S L4 AND (ALLELE? OR POLYMORPHISM? OR MUTATION?)

FILE 'STNGUIDE' ENTERED AT 11:52:39 ON 25 JUL 2002

L5 ANSWER 21 OF 58 MEDLINE
 AN 2000423816 MEDLINE
 DN 20414884 PubMed ID: 10686565
 TI Lack of association between **serotonin transporter**
gene promoter variants and autistic disorder in two ethnically
 distinct samples.
 AU Persico A M; Militeri R; Bravaccio C; Schneider C; Melmed R; Conciatori
 M; Damiani V; Baldi A; Keller F
 CS Laboratory of Neuroscience, Department of Physiology and Neuroscience,
 Libera Universita' "Campus Bio-Medico," Rome, Italy.
 SO AMERICAN JOURNAL OF MEDICAL GENETICS, (2000 Feb 7) 96 (1) 123-7.
 Journal code: 7708900. ISSN: 0148-7299.
 CY United States
 DT Journal; Article; (JOURNAL ARTICLE)
 LA English
 FS Priority Journals
 EM 200009
 ED Entered STN: 20000915
 Last Updated on STN: 20000915
 Entered Medline: 20000907
 AB Family-based studies performed to date provide conflicting evidence of
 linkage/association between autistic disorder and either the "short" [Cook
 et al., 1997: Mol Psychiatry 2:247-250] or the "long" [Klauck et al.,
 1997: Hum Mol Genet 6:2233-2238] **allele** of a polymorphic repeat
 located in the serotonin transporter (5-HTT) gene promoter region,
 affecting 5-HTT gene expression [Lesch et al., 1996: Science
 274:1527-1531]. The present study was designed to assess linkage and
 linkage disequilibrium in two new ethnically distinct samples of families
 with primary autistic probands. The 5-HTT promoter repeat was genotyped in
 54 singleton families collected in Italy and in 32 singleton and 5
 multiplex families collected in the U.S.A., yielding a total sample of 98
 trios. Linkage/association between 5-HTT gene promoter **alleles**
 and autistic disorder was assessed using the transmission/disequilibrium
 test (TDT) and the haplotype-based haplotype relative risk (HHRR). Both
 the Italian and the American samples, either singly or combined, displayed
 no evidence of linkage/association between 5-HTT gene promoter
alleles and autistic disorder. Our findings do not support
 prominent contributions of 5-HTT gene variants to the pathogenesis of
 idiopathic infantile autism. Heterogeneity in pathogenetic mechanisms
 underlying the **disease** may require that linkage/association
 studies be targeted toward patient subgroups isolated on the basis of
 specific biochemical markers, such as serotonin (5-HT) blood levels. Am.
 J. Med. Genet. (Neuropsychiatr. Genet.) 96:123-127, 2000.
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s review and (stress? (4a) (induc?) (4a) disease?))
UNMATCHED RIGHT PARENTHESIS 'DISEASE?))'
The number of right parentheses in a query must be equal to the
number of left parentheses.

=> s review and (stress? (4a) (induc?) (4a) (disease?))
<-----User Break----->